

Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area

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-001

Row 1	Reporter Name [REDACTED]	Submission date.	Contact person (if different than reporter)	Internal ID 2217523
Administrative Data	Address MS USA		Address	
	Phone # [REDACTED]		Phone #	
	Incident Status: New	Location and date of incident MS USA 04/06/2018	Date registrant became aware of incident. 04/15/2018	Was incident part of larger study? No
Row 2	EPA Registration # (Product 1) 53883-59	EPA Registration # (Product 2)	EPA Registration # (Product 3)	
Pesticide(s) Involved	A.I. (s) Glyphosate	A.I. (s)	A.I. (s)	
	Product 1 name Surrender Eraser Systemic Weed & Grass Killer	Product 2 Name	Product 3 Name	
	Exposed to concentrate prior to dilution? No	Exposed to concentrate prior to dilution?	Exposed to concentrate prior to dilution?	
	Formulation Liquid	Formulation	Formulation	
Row 3	Evidence label directions were not followed? No Intentional misuse? No	Incident site: (examples include home, yard, school, industrial, nursery/greenhouse, surface water, commercial turf, building/office, forest/woods, agricultural (specify crop) right-of-way (rail, utility, highway)). Own Residence	Situation (act of using product): (examples include mixing/loading, reentry, application, transportation, repair/ maintenance of application equipment, manufacturing/formulating). See Incident Description Notes	
Applicator certified? UNK				
Incident Circumstances	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff) See Incident Description Notes			

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Brief description of incident circumstances.

Apr 15 2018 9:26AM

Hx: Caller's husband sprayed with the product on 4/6. On the morning of 4/7 he got up to find that he was very dizzy and weak. She took him into an immediate care facility. They took a CXR, found 'some thickening' of his lungs, and gave him a steroid and abx injection. He was dx with walking pneumonia. He now has numbness in his legs, is very dizzy and weak. She also states that the doctors said it was 'sinus' related, but does not elaborate. Is this from the product?

A: Doubtful that husband's sx's are related, but can assist further once we have more product information. Caller becomes very upset when repeatedly asked for and told that we need exact product information (EPA reg #, AI) to be able to properly identify the product and assist her. States we do not know what we are doing and abruptly disconnects.

Apr 16 2018 2:32PM

Client notified

Demographic information: Age: 64 Year(s) Sex: Male Occupation (if relevant) Not specified	Exposure route: Unknown route	Was adverse effect result of suicide/homicide or attempted suicide/homicide? No	Was protective clothing worn (specify)? None Reported
If female, pregnant? NA	Was exposure occupational? Not specified If yes, days lost due to illness: Not specified	Time between exposure and onset of symptoms: Unable to determine	
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician, PCC, hospital inpatient). ER, Urgent Care or Emergent DVM	List signs/symptoms/adverse effects Neurological-Numbness Neurological-Dizziness/vertigo Neurological-Muscle weakness Respiratory-'Thickening' of lungs Respiratory-Walking pneumonia		If lab tests were performed, list test names and results (If available, submit reports) None Reported
Exposure data: NA Amount of pesticide: NA Exposure duration: Acute < 8hrs Patient weight: Unknown			
Human severity category: HC			

This box can be used to provide any explanatory or qualifying information surrounding the incident. (add additional pages if necessary)

Internal ID #
2217523